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| (21) International Application Number: PCT/US99/15308 (22) International Filing Date: 7 July 1999 (07.07.99) (30) Priority Data: 60/092,033 7 July 1998 (07.07.98) US (63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Application US 60/092,033 (CIP) Filed on 7 July 1998 (07.07.98) (71) Applicant (for all designated States except US): THE TRUSTEES OF THE UNIVERSITY OF PENNSYLVANIA [US/US]; Center for Technology Transfer, Suite 300, 3700 Market Street, Philadelphia, PA 19107 (US). (72) Inventor; and (75) Inventor/Applicant (for US only): LU, Zhe [CN/US]; 150 E. Wynnewood #22E, Wynnewood, PA 19096 (US). (74) Agents: LICATA, Jane, Massey et al.; Law Offices of Jane Massey Licata, 66 E. Main Street, Marlton, NJ 08053 (US). | | (81) Designated States: AU, CA, JP, US, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i> |
| (54) Title: COMPOSITIONS AND METHODS FOR INHIBITING INWARD-RECTIFIER POTASSIUM CHANNELS (57) Abstract The present invention provides compounds and methods of identifying and designing compounds which inhibit activity of inward-rectifier K ⁺ channels. In particular, compounds having a tertiapin-like α helix, such as a stable tertiapin derivative wherein the methionine residue in position 13 of tertiapin is replaced by glutamine, are described. Methods of using these compounds to control insulin secretion, and cardiac rhythm and electrical conduction, to modulate neurotransmissions of neurons, and to induce diuresis in mammals are also provided. | | |